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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/506,956	03/24/2005	C. Mauli Agrawal	5660-00503	8795
35690 7590 07/02/2007 MEYERTONS, HOOD, KIVLIN, KOWERT & GOETZEL, P.C. P.O. BOX 398			EXAMINER	
			NAFF, DAVID M	
AUSTIN, TX	/8767-0398		ART UNIT PAPER NUMBER	
			1657	, was 1.
			MAIL DATE	DELIVERY MODE
			07/02/2007	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)				
· · · · · · · · · · · · · · · · · · ·	10/506,956	AGRAWAL ET AL.				
Office Action Summary	Examiner	Art Unit				
	David M. Naff	1657				
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the o	correspondence address				
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DATE of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication.  If NO period for reply is specified above, the maximum statutory period of Failure to reply within the set or extended period for reply will, by statute Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tinuity will apply and will expire SIX (6) MONTHS from, cause the application to become ABANDONE	N. mely filed the mailing date of this communication. ED (35 U.S.C. § 133).				
Status						
1) Responsive to communication(s) filed on 24 M	<u>larch 2005</u> .					
2a) This action is <b>FINAL</b> . 2b) ☐ This	This action is <b>FINAL</b> . 2b)⊠ This action is non-final.					
•	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
closed in accordance with the practice under E	Ex parte Quayle, 1935 C.D. 11, 4	53 O.G. 213.				
Disposition of Claims		•				
4)  Claim(s) 1-32,63 and 128 is/are pending in the 4a) Of the above claim(s) is/are withdraw 5)  Claim(s) is/are allowed.  6)  Claim(s) 1-32,63 and 128 is/are rejected.  7)  Claim(s) is/are objected to.  8)  Claim(s) are subject to restriction and/o	wn from consideration.					
Application Papers	·					
9) The specification is objected to by the Examine 10) The drawing(s) filed on <u>08 September 2004</u> is/a Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex	are: a)⊠ accepted or b)⊡ object drawing(s) be held in abeyance. Se tion is required if the drawing(s) is ob	ee 37 CFR 1.85(a). ojected to. See 37 CFR 1.121(d).				
Priority under 35 U.S.C. § 119						
<ul> <li>12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).</li> <li>a) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents have been received.</li> <li>2. Certified copies of the priority documents have been received in Application No.</li> <li>3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).</li> <li>* See the attached detailed Office action for a list of the certified copies not received.</li> </ul>						
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO/SB/08)  Paper No(s)/Mail Date 3/24/05.	4) ☐ Interview Summar Paper No(s)/Mail [ 5) ☐ Notice of Informal 6) ☐ Other:	Date				

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#### DETAILED ACTION

A preliminary amendment of 9/8/04 canceled claims 33-62, 64-127 and 129.

Claims examined on the merits are 1-32, 63 and 128, which are all claims in the application.

# Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 1-19, 21, 23, 25, 27, 29, 30, 32, 63 and 128 are rejected under 35 U.S.C. 103(a) as being unpatentable over Williams et al (4,927,676) in view of Mineau-Hanschke (6,582,391), and if necessary

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in further view of Hoffman et al (5,034,265) or Lee et al (6,033,582) or Hoffman et al (5,055,316).

The claims are drawn to method of preparing an implant by subjecting a substrate to gas-plasma treatment and exposing the treated substrate to living cells so a portion of the cells become coupled to the substrate, and produce more of a product than cells coupled to an untreated substrate.

Williams et al disclose attaching endothelial cells to a substrate by treating the substrate with a gas-plasma before attaching the cells (col 2, under "SUMMARY OF THE INVENTION", and col 4, lines 48-59).) The substrate with the attached cells is implanted.

Mineau-Hanschke discloses providing a medically useful polypeptide to a patient by providing a matrix containing cells that secrete the polypeptide and implanting the matrix (col 4, lines 17-31). The cells in the matrix can be implanted to produce a wide range of cellular products including various growth factors (col 18, lines 31-55).

Hoffman et al ('265) disclose using gas-plasma treatment to improve compatibility of biomaterials.

Lee et al disclose surface modification of medical implants by gas-plasma treatement.

Hoffman et al ('316) disclose gas-plasma treatment of a surface to provide tight binding of proteins to the surface.

It would have been obvious to use as cells attached to the substrate to be implanted of Williams et al, cells that produce a

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cellular product when implanted disclosed by Mineau-Hanschke to obtain the benefit of cells producing a product in vivo as disclosed by Mineau-Hanschke. Hoffman et al ('265 and '316) and Lee et al further disclose gas-plasma treatment of a substrate, and if needed would have suggested conditions that can be used. Cells attached to the plasma treated substrate will inherently produce more product than cells attached to an untreated substrate. The conditions of dependent claims would have been obvious from conditions disclosed by the references.

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# Claim Rejections - 35 USC § 103

Claim 20 is rejected under 35 U.S.C. 103(a) as being unpatentable over the references as applied to claims 1-19, 21, 23, 25, 27, 29, 30, 32, 63 and 128 above, and further in view of Berlowitz-Tarrant et al (5,840,387).

The claim requires human aortic endothelial cells.

Berlowitz-Tarrant et al disclose attaching aortic endothelial cells to a surface that can be an implant (col 5, lines 40-65).

When using cells that produce a product as the cells of Williams et al as set forth above, it would have been obvious to use aortic endothelial cells as the cells as suggested by Berlowitz-Tarrant et al disclosing attaching aortic endothelial cells to a surface for implanting.

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#### Claim Rejections - 35 USC § 103

Claim 22 is rejected under 35 U.S.C. 103(a) as being unpatentable over the references as applied to claims 1-19, 21, 23, 25, 27, 29, 30, 32, 63 and 128 above, and further in view of Smith et al (5,580,779).

The claim requires myocardial cells.

Smith et al disclose using myocardial cells to produce a peptide in vivo (col 5, lines 3-8).

When using cells that produce a product as the cells of Williams et al as set forth above, it would have been obvious to use myocardial cells to produce a peptide as suggested by Smith et al.

## Claim Rejections - 35 USC § 103

Claim 24 is rejected under 35 U.S.C. 103(a) as being unpatentable over the references as applied to claims 1-19, 21, 23, 25, 27, 29, 30, 32, 63 and 128 above, and further in view of Zonneveld et al (6,447,768).

The claim requires the cellular product to be a nucleic acid.

Zonneveld et al disclose delivering a nucleic acid *in vivo* with a cell that produces the nucleic acid to provide gene therapy (abstract and paragraph bridging cols 3 and 4.

When using cells that produce a product as the cells of
Williams et al as set forth above, it would have been obvious to use
cells that produce a nucleic acid to provide gene therapy as suggested
by Zonneveld et al.

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#### Claim Rejections - 35 USC § 103

Claim 26 is rejected under 35 U.S.C. 103(a) as being unpatentable over the references as applied to claims 1-19, 21, 23, 25, 27, 29, 30, 32, 63 and 128 above, and further in view of Beckmann et al (6,306,615).

The claim requires the cellular product to be beta-tubulin.

Beckmann et al disclose beta-tubulin-producing cells (col 16, lines 15-19).

When using cells that produce a product as the cells of Williams

10 et al as set forth above, it would have been obvious to use cells that

produce beta-tubulin to obtain its function as suggested by Beckmann

et al.

# Claim Rejections - 35 USC § 103

Claim 28 is rejected under 35 U.S.C. 103(a) as being unpatentable

over the references as applied to claims 1-19, 21, 23, 25, 27, 29, 30,

32, 63 and 128 above, and further in view of Mineau-Hanschke

(6,419,920).

The claim requires vascular endothelial growth factor as the cellular product.

20 Mineau-Hanschke discloses vascular endothelial growth factor as a cellular product (col 4, lines 41-42).

When using cells that produce a product as the cells of Williams et al as set forth above, it would have been obvious to use cells that produce vascular endothelial growth factor as suggested by Mineau-

25 Hanschke.

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### Claim Rejections - 35 USC § 103

Claim 31 is rejected under 35 U.S.C. 103(a) as being unpatentable over the references as applied to claims 1-19, 21, 23, 25, 27, 29, 30, 32, 63 and 128 above, and further in view of Newman et al (6,087,331).

The claim requires platelet-endothelial cell adhesion molecule-1 as a cellular product.

Newman et al disclose therapeutic use of platelet-endothelial cell adhesion molecule-1 and cells transformed to produce platelet-endothelial cell adhesion molecule-1.

When using cells that produce a product as the cells of Williams et al as set forth above, it would have been obvious to use cells that produce platelet-endothelial cell adhesion molecule-1 as suggested by Newman et al.

#### Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to David M. Naff whose telephone number is 571-272-0920. The examiner can normally be reached on Monday-Friday 9:30-6:00.

If attempts to reach the examiner by telephone are unsuccessful,

the examiner's supervisor, Jon Weber can be reached on 571-272-0925.

The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

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